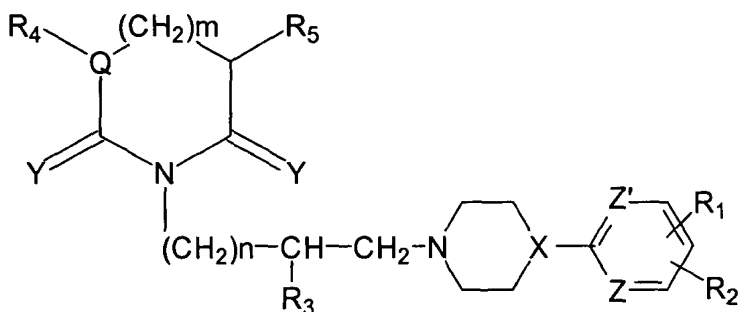


In the claims:

Kindly amend claims 44 and 45 to read as follows:

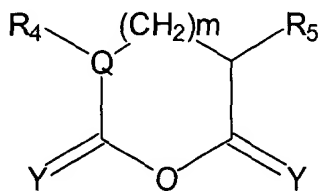
44. (Four Times Amended) A method for making a compound having the structure of Formula I

El



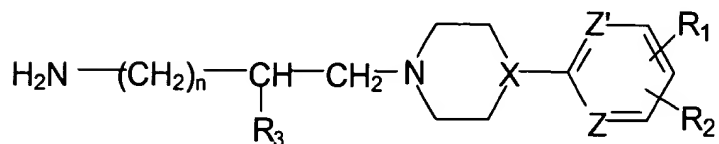
(I)

its pharmaceutically acceptable salts, enantiomers, diastereomers, or N-oxides, wherein Y is O or S; Q, Z and Z' are independently CH; X is CH or N; m=0; n=0-4; R₁, R₂ are independently selected from: F, Br, OCH₃, OC₂H₅, OCH₂CF₃, SCF₃, C₂H₅, CF₃, isopropoxy, and cyclopropyl; and R₃, R₄ and R₅ are independently H, C₁₋₃ alkyl, substituted or unsubstituted phenyl, which comprises reacting a compound having the structure of Formula VI'



(VI')

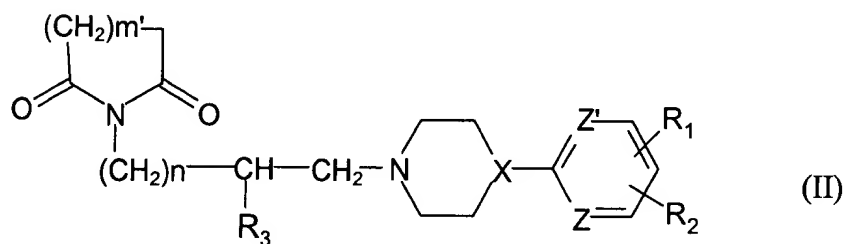
with a compound having the structure of Formula V in pyridine at reflux temperature followed by reflux in the presence of acetic anhydride



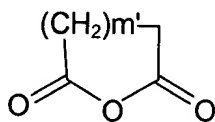
(V)

thereby to produce the compound of Formula I.

45. (Thrice Amended) A method for making a compound having the structure of Formula II

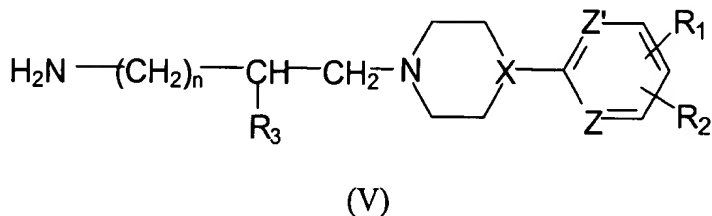


its pharmaceutically acceptable salts, enantiomers, diastereomers, or N-oxides, wherein X is CH or N; Z and Z' are independently CH; n = 0-4; m' = 1; R₁, R₂ are independently selected from: F, Br, OCH₃, OC₂H₅, OCH₂CF₃, SCF₃, isopropoxy, and cyclopropyl; and R₃ is independently H, C₁₋₃ alkyl, substituted or unsubstituted phenyl, which comprises reacting a compound having the structure of Formula VI



(VI)

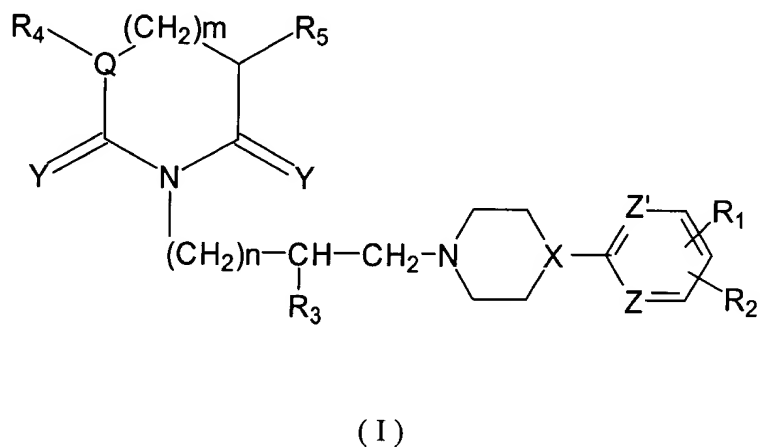
with a compound having the structure of Formula V in pyridine at reflux temperature followed by reflux in the presence of acetic anhydride



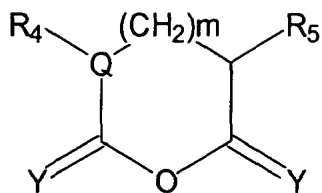
thereby to produce the compound of Formula II.

Please add new claims 48 and 49.

48. (New) A method for making a compound having the structure of Formula I

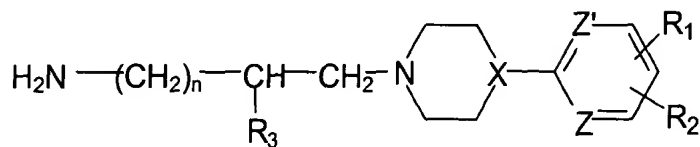


its pharmaceutically acceptable salts, enantiomers, diastereomers, or N-oxides, wherein Y is O or S; Q, Z and Z' are independently CH; X is CH or N; m=1-3; n=0-4; R₁, R₂ are independently selected from: F, Cl, Br, OC₂H₅, OCH₂CF₃, SCF₃, CH₃, C₂H₅, CF₃, isopropoxy, and cyclopropyl; and R₃, R₄ and R₅ are independently H, C₁₋₃ alkyl, substituted or unsubstituted phenyl, which comprises reacting a compound having the structure of Formula VI'



(VI')

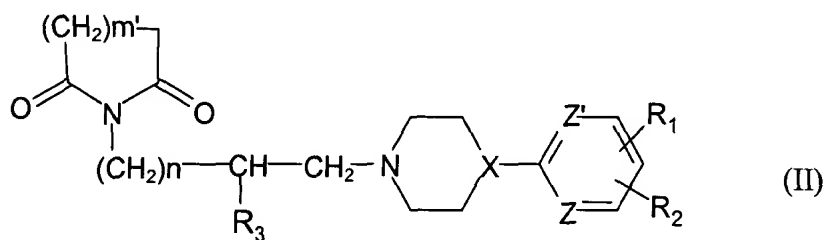
with a compound having the structure of Formula V in pyridine at reflux temperature followed by reflux in the presence of acetic anhydride



(V)

thereby to produce the compound of Formula I.

49. (New) A method for making a compound having the structure of Formula II



(II)

its pharmaceutically acceptable salts, enantiomers, diastereomers, or N-oxides, wherein X is CH or N; Z and Z' are independently CH; n = 0-4; m' = 2-4; R₁, R₂ are independently selected from: F, Cl, Br, OC₂H₅, OCH₂CF₃, SCF₃, CH₃, isopropoxy, and cyclopropyl; and R₃ is independently H, C₁₋₃ alkyl, substituted or unsubstituted phenyl, which comprises reacting a compound having the structure of Formula VI